## Amendments to the claims

This listing of claims will replace all prior versions, and listings, of claims in the application:

1. (Currently amended): A compound of formula (I)

wherein:

Z<sub>1</sub> is N;

R1 and R1a\_are independently is\_hydrogen; hydroxy;  $(C_{1-6})$ alkoxy unsubstituted or substituted by  $(C_{1-6})$ alkoxy, hydroxy, amino, piperidyl, guanidino or amidino any of which is unsubstituted or N-substituted by one or two  $(C_{1-6})$ alkyl, acyl,  $(C_{1-6})$ alkylsulphonyl, CONH2, hydroxy,  $(C_{1-6})$ alkylthio, heterocyclylthio, heterocyclyloxy, arylthio, aryloxy, acylthio, acyloxy or  $(C_{1-6})$ alkylsulphonyloxy;  $(C_{1-6})$ alkoxysubstituted $(C_{1-6})$ alkyl; halogen;  $(C_{1-6})$ alkyl;  $(C_{1-6})$ alkylthio; trifluoromethyl; trifluoromethoxy; nitro; azido; cyano; acyl; acyloxy; acylthio;  $(C_{1-6})$ alkylsulphonyl;  $(C_{1-6})$ alkylsulphoxide; arylsulphonyl; arylsulphoxide; or an amino, piperidyl, guanidino or amidino group unsubstituted or N-substituted by one or two  $(C_{1-6})$ alkyl, acyl or  $(C_{1-6})$ alkylsulphonyl groups; or R1-and R1a-may together form ethylenedioxy;

 ${\sf R}^2$  is H or halogen;

provided that when  $Z_1$  is N, then  $R^2$  is H;

R³ is hydrogen; halogen; hydroxy; cyano; CF₃; nitro; azido; acyl; aryl; heteroaryl;  $CO_2H$ ; aeyexyacyloxy; acylthio;  $(C_{1-6})$ alkyl unsubstituted or substituted by one or two  $(C_{1-6})$ alkoxy, hydroxy, amino, piperidyl, guanidino or amidino any of which is unsubstituted or N-substituted by one or two  $(C_{1-6})$ alkyl, acyl,  $(C_{1-6})$ alkylsulphonyl,  $CONH_2$ , hydroxy,  $(C_{1-6})$ alkylthio, heterocyclylthio, heterocyclyloxy, arylthio, aryloxy, acylthio, acyloxy or  $(C_{1-6})$ alkylsulphonyloxy;  $(C_{1-6})$ alkoxy unsubstituted or substituted by one or two  $(C_{1-6})$ alkoxy, hydroxy, amino, piperidyl, guanidino or amidino any of which is unsubstituted or N-substituted by one or two  $(C_{1-6})$ alkyl, acyl,  $(C_{1-6})$ alkylsulphonyl,  $CONH_2$ , hydroxy,  $(C_{1-6})$ alkylthio, heterocyclylthio, heterocyclyloxy, arylthio, aryloxy, acylthio, acyloxy or  $(C_{1-6})$ alkylsulphonyloxy;  $(C_{3-7})$ cycloalkyl;  $(C_{1-6})$ alkoxy-substituted $(C_{1-6})$ alkyl;  $(C_{1-6})$ alkylsulphonyl; or arylsulphoxide; or an amino, piperidyl, guanidino or amidino group unsubstituted or N-substituted by one or two  $(C_{1-6})$ alkyl, acyl or  $(C_{1-6})$ alkylsulphonyl groups;

```
w_1 is N, C, or CR^4;

w_2 is C=O, CR^4, or CR^4R^5;

w_3 is C=O or CR^4R^5;

w_4 is N or CR^4;

w_5 is C=O or CR^4R^5;

w_6 is C=O, CR^4, or CR^4R^5;
```

or, one of W<sub>2</sub>, W<sub>3</sub>, W<sub>5</sub> and W<sub>6</sub> is CR<sup>4</sup>R<sup>5</sup>CR<sup>4</sup>R<sup>5</sup> and the others are defined as above; wherein each R<sup>4</sup> and R<sup>5</sup> is independently hydrogen; halogen; hydroxy; cyano; CF<sub>3</sub>; nitro; azido; acyl; aryl; heteroaryl; CO<sub>2</sub>H; acyexyacyloxy; acylthio; (C<sub>1-6</sub>)alkyl unsubstituted or substituted by one or two (C<sub>1-6</sub>)alkoxy, hydroxy, amino, piperidyl, guanidino or amidino any of which is unsubstituted or N-substituted by one or two (C<sub>1-6</sub>)alkyl, acyl, (C<sub>1-6</sub>)alkylsulphonyl, CONH<sub>2</sub>, hydroxy, (C<sub>1-6</sub>)alkylthio, heterocyclylthio, heterocyclyloxy, arylthio, aryloxy, acylthio, acyloxy or (C<sub>1-6</sub>)

6)alkylsulphonyloxy; ( $C_{1-6}$ )alkoxy unsubstituted or substituted by one or two ( $C_{1-6}$ )alkoxy, hydroxy, amino, piperidyl, guanidino or amidino any of which is unsubstituted or N-substituted by one or two ( $C_{1-6}$ )alkyl, acyl, ( $C_{1-6}$ )alkylsulphonyl, CONH<sub>2</sub>, hydroxy, ( $C_{1-6}$ )alkylthio, heterocyclylthio, heterocyclyloxy, arylthio, aryloxy, acylthio, acyloxy or ( $C_{1-6}$ )alkylsulphonyloxy; ( $C_{3-7}$ )cycloalkyl; ( $C_{1-6}$ )alkoxy-substituted( $C_{1-6}$ )alkyl; ( $C_{1-6}$ )alkylthio; trifluoromethoxy; ( $C_{1-6}$ )alkylsulphonyl; ( $C_{1-6}$ )alkylsulphoxide; arylsulphonyl; or arylsulphoxide; or an amino, piperidyl, guanidino or amidino group unsubstituted or N-substituted by one or two ( $C_{1-6}$ )alkyl, acyl or ( $C_{1-6}$ )alkylsulphonyl groups; or two R<sup>5</sup> groups are joined together to form bicycloheptane;

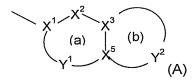
A is  $CR^6R^7$  or C(O); B is  $CR^8R^9$  or C(O);

wherein R<sup>6</sup>, R<sup>7</sup>, R<sup>8</sup>, and R<sup>9</sup> are independently hydrogen; halogen; hydroxy; cyano; CF<sub>3</sub>; nitro; azido; acyl; aryl; heteroaryl; CO<sub>2</sub>H; aeyexyacyloxy; acylthio; (C<sub>1-6</sub>)alkyl unsubstituted or substituted by one or two (C<sub>1-6</sub>)alkoxy, hydroxy, amino, piperidyl, guanidino or amidino any of which is unsubstituted or N-substituted by one or two (C<sub>1-6</sub>)alkyl, acyl, (C<sub>1-6</sub>)alkylsulphonyl, CONH<sub>2</sub>, hydroxy, (C<sub>1-6</sub>)alkylthio, heterocyclylthio, heterocyclyloxy, arylthio, aryloxy, acylthio, acyloxy or (C<sub>1-6</sub>)alkylsulphonyloxy; (C<sub>1-6</sub>)alkoxy unsubstituted or substituted by one or two (C<sub>1-6</sub>)alkoxy, hydroxy, amino, piperidyl, guanidino or amidino any of which is unsubstituted or N-substituted by one or two (C<sub>1-6</sub>)alkyl, acyl, (C<sub>1-6</sub>)alkylsulphonyl, CONH<sub>2</sub>, hydroxy, (C<sub>1-6</sub>)alkylthio, heterocyclylthio, heterocyclyloxy, arylthio, aryloxy, acylthio, acyloxy or (C<sub>1-6</sub>)alkylsulphonyloxy; (C<sub>3-7</sub>)cycloalkyl; (C<sub>1-6</sub>)alkoxy-substituted(C<sub>1-6</sub>)alkyl; (C<sub>1-6</sub>)alkylthio; trifluoromethoxy; (C<sub>1-6</sub>)alkylsulphonyl; (C<sub>1-6</sub>)alkylsulphonyl; or arylsulphoxide; or an amino, piperidyl, guanidino or amidino

group unsubstituted or N-substituted by one or two ( $C_{1-6}$ )alkyl, acyl or ( $C_{1-6}$ )alkylsulphonyl groups;

R<sup>10</sup> is hydrogen; aryl; heteroaryl; (C<sub>1-6</sub>)alkyl unsubstituted or substituted by one or two (C<sub>1-6</sub>)alkoxy, <u>acyloxy, carboxy,</u> hydroxy, amino, piperidyl, piperazinyl, morpholino, guanidino, or amidino, any of which is unsubstituted or N-substituted by one or two aryl, heteroaryl, halogen, cyano, CF<sub>3</sub>, unsubstituted (C<sub>1-6</sub>)alkyl, acyl, (C<sub>1-6</sub>)alkylsulphonyl, arylsulphonyl, hydroxy, (C<sub>1-6</sub>)alkylthio, heterocyclylthio, heterocyclyloxy, arylthio, aryloxy, acylthio, acyloxy, or (C<sub>1-6</sub>)alkylsulphonyloxy, provided that the substitution does not lead to an unstable compound; (C<sub>1-6</sub>)alkoxy-substituted(C<sub>1-6</sub>)alkyl; hydroxy-substituted(C<sub>1-6</sub>)alkyl; (C<sub>1-6</sub>)alkylcarbonyl; (C<sub>2-6</sub>)alkenylcarbonyl; (C<sub>1-6</sub>)alkoxycarbonyl; CO<sub>2</sub>H; or CF<sub>3</sub>;

R<sup>11</sup> is a group -U-R<sup>12</sup> where R<sup>12</sup> is a substituted or unsubstituted bicyclic carbocyclic or heterocyclic ring system (A):



containing up to four heteroatoms in each ring in which at least one of rings (a) and (b) is aromatic;

 $\mathsf{X}^1$  is C or N when part of an aromatic ring or  $\mathsf{CR}^{14}$  when part of a non aromatic ring;

 $X^2$  is N, NR<sup>13</sup>, O, S(O)<sub>X</sub>, CO or CR<sup>14</sup> when part of an aromatic or non-aromatic ring or may in addition be CR<sup>14</sup>R<sup>15</sup> when part of a non aromatic ring;

 $X^3$  and  $X^5$  are independently N or C;

 $Y^1$  is a 0 to 4 atom linker group each atom of which is independently selected from N, NR<sup>13</sup>, O, S(O)<sub>X</sub>, CO and CR<sup>14</sup> when part of an aromatic or non-aromatic ring or may additionally be CR<sup>14</sup>R<sup>15</sup> when part of a non aromatic ring,

 $Y^2$  is a 2 to 6 atom linker group, each atom of  $Y^2$  being independently selected from N, NR<sup>13</sup>, O, S(O)<sub>X</sub>, CO and CR<sup>14</sup> when part of an aromatic or non-aromatic ring or may additionally be CR<sup>14</sup>R<sup>15</sup> when part of a non aromatic ring;

each of R<sup>14</sup> and R<sup>15</sup> is independently selected from: H; (C<sub>1-4</sub>)alkylthio; halo; (C<sub>1-4</sub>)alkyl; (C<sub>2-4</sub>)alkenyl; hydroxy; hydroxy(C<sub>1-4</sub>)alkyl; mercapto(C<sub>1-4</sub>)alkyl; (C<sub>1-4</sub>)alkoxy; trifluoromethoxy; nitro; cyano; carboxy; amino or aminocarbonyl unsubstituted or substituted by (C<sub>1-4</sub>)alkyl;

each  $R^{13}$  is independently H; trifluoromethyl;  $(C_{1-4})$ alkyl unsubstituted or substituted by hydroxy, carboxy,  $(C_{1-4})$ alkoxy,  $(C_{1-6})$ alkylthio, halo or trifluoromethyl;  $(C_{2-4})$ alkenyl; or aminocarbonyl wherein the amino group is optionally substituted  $(C_{1-4})$ alkyl;

each x is independently 0, 1 or 2; U is CO, SO<sub>2</sub>, CH<sub>2</sub>, or CR<sup>16</sup>R<sup>17</sup>;

R<sup>16</sup> and R<sup>17</sup> are independently selected from H; aryl; heteroaryl; (C<sub>1-6</sub>)alkyl; (C<sub>1-6</sub>)alkyl substituted by (C<sub>1-6</sub>)alkoxy, hydroxy, amino, piperidyl, piperazinyl, morpholino, guanidino, or amidino, any of which is substituted or N-substituted by one or two H, aryl, heteroaryl, halogen, cyano, CF<sub>3</sub>, (C<sub>1-6</sub>)alkyl, acyl, (C<sub>1-6</sub>)alkylsulphonyl, arylsulphonyl, hydroxy, (C<sub>1-6</sub>)alkylthio, heterocyclyloxy, arylthio, aryloxy, acylthio, acyloxy, or (C<sub>1-6</sub>)alkylsulphonyloxy, provided that the substitution does not lead to an unstable compound; (C<sub>1-6</sub>)alkoxy-substituted(C<sub>1-6</sub>)alkyl; hydroxy-substituted(C<sub>1-6</sub>)alkyl; amino-substituted(C<sub>1-6</sub>)alkyl, which is N-substituted by one or two (C<sub>1-6</sub>)alkyl, acyl, (C<sub>1-6</sub>)alkylsulphonyl, or arylsulphonyl; (C<sub>1-6</sub>)alkylcarbonyl; (C<sub>2-6</sub>)alkenylcarbonyl;

 $(C_{1-6})$ alkoxycarbonyl;  $CO_2H$ ; or  $CF_3$ ; or

a pharmaceutically acceptable salt thereof.

2. (Previously presented): A compound or salt according to claim 1, wherein  $R^1$  is F, Cl, OCH<sub>3</sub>, methyl, or SCH<sub>3</sub>.

- 3. Canceled.
- 4. (Previously presented): A compound or salt according to claim 1, wherein  $\mathbb{R}^2$  is H or F.
- 5. (Previously presented): A compound or salt according to claim 1, wherein  $\mathbb{R}^3$  is CI or F.
- 6. (Previously presented): A compound or salt according to claim 1, wherein each R<sup>4</sup> is independently H, methyl, OH, COOH, NH<sub>2</sub>, OCH<sub>3</sub>, or CH<sub>2</sub>OH.
- 7. (Previously presented): A compound or salt according to claim 1, wherein  $\mathbb{R}^5$  is H.
- 8. (Previously presented): A compound or salt according to claim 1, wherein the group –U– is –CH<sub>2</sub>–.
- 9. (Previously presented): A compound or salt according to claim 1, wherein R<sup>12</sup> is: benzo[1,2,5]thiadiazol-5-yl;

4H-benzo[1,4] thiazin-3-one-6-yl;

2,3-dihydro-benzo[1,4]dioxin-6-yl;

benzo[1,2,3]thiadiazol-5-yl;

3-oxo-3,4-dihydro-2H-benzo[1,4]oxazin-6-yl;

7-fluoro-3-oxo-3,4-dihydro-2H-benzo[1,4] oxazin-6-yl;

2-oxo-2,3-dihydro-1H-pyrido[2,3-b][1,4]thiazin-7-yl;

2,3-Dihydro-[1,4]dioxino[2,3-c]pyridin-7-yl;

3-oxo-3,4-dihydro-2H-pyrido[3,2-b][1,4]oxazin-6-yl;

[1,2,3]thiadiazolo[5,4-b]pyridin-6-yl;

3-oxo-3,4-dihydro-2*H*-pyrido[3,2-*b*][1,4]thiazin-6-yl;

7-chloro-3-oxo-3,4-dihydro-2*H*-pyrido[3,2-*b*][1,4]thiazin-6-yl;

7-fluoro-3-oxo-3,4-dihydro-2H-pyrido[3,2-b][1,4]thiazin-6-yl; or

2-oxo-2,3-dihydro-1*H*-pyrido[3,4-*b*][1,4]thiazin-7-yl.

10. (Currently amended): A compound according to claim 1, wherein the compound is:

6-({2-[1-(6-methoxynaphthyridin-4-yl)piperidin-4-yl]ethylamino} methyl)-4*H*-pyrido[3,2-*b*][1,4]oxazin-3-one;

6-({2-[1-(6-methoxynaphthyridin-4-yl)piperidin-4-yl]ethylamino} methyl)-4*H*-pyrido[3,2-*b*][1,4]thiazin-3-one;

(2,3-dihydro-[1,4]dioxino[2,3-c]pyridin-7-ylmethyl)-{2-[1-(6-methoxynaphthyridin-4-yl)piperidin-4-yl]ethyl}amine;

6-({2-[1-(3-chloro-6-methoxy-[1,5]naphthyridin-4-yl)phenyl] ethylamino}methyl)-4*H*-pyrido[3,2-*b*][1,4]oxazin-3-one;

6-({2-[1-(3-chloro-6-methoxy-[1,5]naphthyridin-4-yl)phenyl] ethylamino}methyl)-4*H*-pyrido[3,2-*b*][1,4]thiazin-3-one;

 $\{2-[1-(3-chloro-6-methoxynaphthyridin-4-yl)piperidin-4-yl]ethyl\}-(2,3-dihydro[1,4]dioxino[2,3-c]pyridin-7-ylmethyl)amine;$ 

 $6-(\{2-[4-(6-methoxynaphthyridin-4-yl)piperazin-1-yl]$  ethylamino} methyl)-4H-pyrido[3,2-b][1,4]oxazin-3-one;

6-({2-[4-(6-methoxynaphthyridin-4-yl)piperazin-1-yl] ethylamino} methyl)-4*H*-pyrido[3,2-*b*][1,4]thiazin-3-one;

(2,3-dihydro-[1,4]dioxino[2,3-c]pyridin-7-ylmethyl)-{2-[4-(6-methoxynaphthyridin-4-yl)piperizin-1-yl]ethyl}amine;

6-({2-[4-(3-chloro-6-methoxynaphthyridin-4-yl)piperazin-1-yl] ethylamino}methyl)-4*H*-pyrido[3,2-*b*][1,4]oxazin-3-one;

6-({2-[4-(3-chloro-6-methoxynaphthyridin-4-yl)piperazin-1-yl] ethylamino}methyl)-4*H*-pyrido[3,2-*b*][1,4]thiazin-3-one;

 $\{2-[4-(3-chloro-6-methoxynaphthyridin-4-yl)piperazin-1-yl]ethyl\}-(2,3-dihydro[1,4]dioxino[2,3-c]pyridin-7-ylmethyl)amine;$ 

6-({2-[4-(6-methoxy-[1,5]naphthyridin-4-yl)-3,6-dihydro-2 H -pyridin-1-yl]-2-oxo-ethylamino}-methyl) -4 H -pyrido[3,2-b][1,4]thiazin-3-one;

N-(2-{1-[6-(methyloxy)-1,5-naphthyridin-4-yl]-4-piperidinyl}ethyl)-3-oxo-3,4-dihydro-2H-pyrido[3,2-b][1,4]thiazine-6-carboxamide;

N-(2-{1-[6-(methyloxy)-1,5-naphthyridin-4-yl]-4-piperidinyl}ethyl)-3-oxo-3,4-dihydro-2H-1,4-benzothiazine-6-sulfonamide;

N-methyl-N-(2-{1-[6-(methyloxy)-1,5-naphthyridin-4-yl]-4-piperidinyl} ethyl)-3-oxo-3,4-dihydro-2H-pyrido[3,2-b][1,4]thiazine-6-carboxamide;

N-methyl-N-(2-{1-[6-(methyloxy)-1,5-naphthyridin-4-yl]-4-piperidinyl} ethyl)-3-oxo-3,4-dihydro-2H-1,4-benzothiazine-6-sulfonamide;

N-(2-{1-[3-chloro-6-(methyloxy)-1,5-naphthyridin-4-yl]-4-piperidinyl} ethyl)-3-oxo-3,4-dihydro-2H-1,4-benzothiazine-6-sulfonamide;

7-{[(2-{4-[6-(methyloxy)-1,5-naphthyridin-4-yl]-1-piperazinyl}ethyl) oxy] methyl}-2,3-dihydro[1,4]dioxino[2,3-c]pyridine;

N-(2-{4-[6-(methyloxy)-1,5-naphthyridin-4-yl]-1-piperazinyl}ethyl)-3-oxo-3,4-dihydro-2H-pyrido[3,2-b][1,4]thiazine-6-carboxamide;

N-methyl-N-(2-{4-[6-(methyloxy)-1,5-naphthyridin-4-yl]-1-piperazinyl}ethyl)-3-oxo-3,4-dihydro-2H-pyrido[3,2-b][1,4]thiazine-6-carboxamide;

N-(2-{4-[6-(methyloxy)-1,5-naphthyridin-4-yl]-1-piperazinyl}ethyl)-3-oxo-3,4-dihydro-2H-1,4-benzothiazine-6-sulfonamide;

N-methyl-N-(2-{4-[6-(methyloxy)-1,5-naphthyridin-4-yl]-1-piperazinyl} ethyl)-3-oxo-3.4-dihydro-2H-1,4-benzothiazine-6-sulfonamide;

6-{[(2-{4-[6-(methyloxy)-1,5-naphthyridin-4-yl]hexahydro-1H-1,4-diazepin-1-yl}ethyl)amino]methyl}-2H-pyrido[3,2-b][1,4]thiazin-3(4H)-one;

N-(2-{4-[6-(methyloxy)-1,5-naphthyridin-4-yl]hexahydro-1H-1,4-diazepin-1-yl}ethyl)-3-oxo-3,4-dihydro-2H-1,4-benzothiazine-6-sulfonamide;

6-{[(2-{(1R,4R)-5-[6-(methyloxy)-1,5-naphthyridin-4-yl]-2,5-diazabicyclo [2.2.1]hept-2-yl}ethyl)amino]methyl}-2H-pyrido[3,2-b][1,4]thiazin-3(4H)-one;

6-[({1-[6-(methyloxy)-1,5-naphthyridin-4-yl]-4-piperidinyl}amino) methyl]-2H-pyrido[3,2-b][1,4]thiazin-3(4H)-one;

6-{[(2-{4-hydroxy-1-[6-(methyloxy)-1,5-naphthyridin-4-yl]-4-piperidinyl}ethyl) amino]methyl}-2H-pyrido[3,2-b][1,4]thiazin-3(4H)-one;

6-{[(2-{4-hydroxy-1-[6-(methyloxy)-1,5-naphthyridin-4-yl]-4-piperidinyl}ethyl) amino|methyl}-2H-pyrido[3,2-b][1,4]oxazin-3(4H)-one;

N-(2-{4-hydroxy-1-[6-(methyloxy)-1,5-naphthyridin-4-yl]-4-piperidinyl} ethyl)-3-oxo-3,4-dihydro-2H-1,4-benzothiazine-6-sulfonamide;

6-{[(2-{4-[7-fluoro-6-(methyloxy)-1,5-naphthyridin-4-yl]-1-piperazinyl} ethyl) amino]methyl}-2H-pyrido[3,2-b][1,4]thiazin-3(4H)-one; or

6-{[(2-{4-[7-fluoro-6-(methyloxy)-1,5-naphthyridin-4-yl]-1-piperazinyl} ethyl)amino]methyl}-2H-pyrido[3,2-b][1,4]oxazin-3(4H)-one; or a pharmaceutically acceptable salt thereof.

11. (Previously presented): A pharmaceutical composition, comprising a compound or salt according to claim 1 and a pharmaceutically acceptable carrier.

- 12. (Previously presented): A method of treating bacterial infections in mammals, which comprises administering to a mammal in need thereof an effective amount of a compound or salt according to claim 1.
- 13. (Previously presented) A method according to claim 12, wherein the mammal is a human.
- 14. (Previously presented) A compound or salt according to claim 1, wherein  $R^{12}$  is an aromatic heterocyclic ring (A) having 8-11 ring atoms including 2-4 heteroatoms of which at least one is N or NR<sup>13</sup>, in which preferably Y<sup>2</sup> contains 2-3 heteroatoms, one of which is S and 1-2 are N, with one N bonded to X<sup>3</sup>.
- 15. (Previously presented) A compound or salt according to claim 1, wherein  $R^{12}$  has a heterocyclic ring (A) having ring (a) aromatic selected from optionally substituted benzo and pyrido and ring (b) non-aromatic and in which  $Y^2$  has 3-5 atoms including a heteroatom bonded to  $X^5$  selected from NR<sup>13</sup>, O or S and NHCO bonded via N to  $X^3$ . or O bonded to  $X^3$ .
- 16. (Previously presented) A compound or salt according to claim 1, wherein R<sup>12</sup> is: 3-oxo-3,4-dihydro-2*H*-pyrido[3,2-*b*][1,4]thiazin-6-yl, 3-oxo-3,4-dihydro-2*H*-pyrido[3,2-*b*][1,4]oxazin-6-yl, or 2,3-dihydro-[1,4]dioxino[2,3-c]pyridin-7-yl.
- 17. (Previously presented) A compound according to claim 1, wherein the compound is:

6-({2-[1-(6-methoxynaphthyridin-4-yl)piperidin-4-yl]ethylamino} methyl)-4*H*-pyrido[3,2-*b*][1,4]oxazin-3-one;

6-({2-[1-(3-chloro-6-methoxy-[1,5]naphthyridin-4-yl)phenyl] ethylamino}methyl)-4*H*-pyrido[3,2-*b*][1,4]oxazin-3-one;

6-({2-[4-(6-methoxynaphthyridin-4-yl)piperazin-1-yl] ethylamino} methyl)-4*H*-pyrido[3,2-*b*][1,4]oxazin-3-one;

6-({2-[4-(3-chloro-6-methoxynaphthyridin-4-yl)piperazin-1-yl] ethylamino}methyl)-4*H*-pyrido[3,2-*b*][1,4]oxazin-3-one;

6-{[(2-{4-hydroxy-1-[6-(methyloxy)-1,5-naphthyridin-4-yl]-4-piperidinyl}ethyl) amino]methyl}-2H-pyrido[3,2-b][1,4]oxazin-3(4H)-one; or

6-{[(2-{4-[7-fluoro-6-(methyloxy)-1,5-naphthyridin-4-yl]-1-piperazinyl}

ethyl)amino]methyl}-2H-pyrido[3,2-b][1,4]oxazin-3(4H)-one; or a pharmaceutically acceptable salt thereof.

18. (Currently amended) A compound according to claim 1, wherein the compound is:

6-({2-[1-(6-methoxynaphthyridin-4-yl)piperidin-4-yl]ethylamino} methyl)-4*H*-pyrido[3,2-*b*][1,4]thiazin-3-one;

6-({2-[1-(3-chloro-6-methoxy-[1,5]naphthyridin-4-yl)phenyl] ethylamino}methyl)-4*H*-pyrido[3,2-*b*][1,4]thiazin-3-one;

6-({2-[4-(6-methoxynaphthyridin-4-yl)piperazin-1-yl] ethylamino} methyl)-4*H*-pyrido[3,2-*b*][1,4]thiazin-3-one;

 $6-(\{2-[4-(3-chloro-6-methoxynaphthyridin-4-yl)piperazin-1-yl] ethylamino\}methyl)-4H-pyrido[3,2-b][1,4]thiazin-3-one;$ 

N-(2-{1-[6-(methyloxy)-1,5-naphthyridin-4-yl]-4-piperidinyl}ethyl)-3-oxo-3,4-dihydro-2H-pyrido[3,2-b][1,4]thiazine-6-carboxamide;

N-methyl-N-(2-{1-[6-(methyloxy)-1,5-naphthyridin-4-yl]-4-piperidinyl} ethyl)-3-oxo-3,4-dihydro-2H-pyrido[3,2-b][1,4]thiazine-6-carboxamide;

N-(2-{4-[6-(methyloxy)-1,5-naphthyridin-4-yl]-1-piperazinyl}ethyl)-3-oxo-3,4-dihydro-2H-pyrido[3,2-b][1,4]thiazine-6-carboxamide;

N-methyl-N-(2-{4-[6-(methyloxy)-1,5-naphthyridin-4-yl]-1-piperazinyl}ethyl)-3-oxo-3,4-dihydro-2H-pyrido[3,2-b][1,4]thiazine-6-carboxamide;

6-{[(2-{4-[6-(methyloxy)-1,5-naphthyridin-4-yl]hexahydro-1H-1,4-diazepin-1-yl}ethyl)amino]methyl}-2H-pyrido[3,2-b][1,4]thiazin-3(4H)-one;

6-{[(2-{(1R,4R)-5-[6-(methyloxy)-1,5-naphthyridin-4-yl]-2,5-diazabicyclo [2.2.1]hept-2-yl}ethyl)amino]methyl}-2H-pyrido[3,2-b][1,4]thiazin-3(4H)-one;

6-[({1-[6-(methyloxy)-1,5-naphthyridin-4-yl]-4-piperidinyl}amino) methyl]-2H-pyrido[3,2-b][1,4]thiazin-3(4H)-one;

 $6-\{[(2-\{4-hydroxy-1-[6-(methyloxy)-1,5-naphthyridin-4-yl]-4-piperidinyl\}ethyl) amino]methyl\}-2H-pyrido[3,2-b][1,4]thiazin-3(4H)-one; or \\$ 

 $6-\{[(2-\{4-[7-fluoro-6-(methyloxy)-1,5-naphthyridin-4-yl]-1-piperazinyl\}\ ethyl)$  amino]methyl}-2H-pyrido[3,2-b][1,4]thiazin-3(4H)-one; or

a pharmaceutically acceptable salt thereof.

19. (Currently amended) A compound according to claim 1, wherein the compound is:

- (2,3-dihydro-[1,4]dioxino[2,3-c]pyridin-7-ylmethyl)-{2-[1-(6-methoxynaphthyridin-4-yl)piperidin-4-yl]ethyl}amine;
- {2-[1-(3-chloro-6-methoxynaphthyridin-4-yl)piperidin-4-yl]ethyl}-(2,3-dihydro[1,4]dioxino[2,3-c]pyridin-7-ylmethyl)amine;
- (2,3-dihydro-[1,4]dioxino[2,3-c]pyridin-7-ylmethyl)-{2-[4-(6-methoxynaphthyridin-4-yl)piperizin-1-yl]ethyl}amine;
- $\{2-[4-(3-chloro-6-methoxynaphthyridin-4-yl)piperazin-1-yl]ethyl\}-(2,3-dihydro[1,4]dioxino[2,3-c]pyridin-7-ylmethyl)amine; or$
- $\frac{7-\{[(2-\{4-[6-(methyloxy)-1,5-naphthyridin-4-yl]-1-piperazinyl\}ethyl)\cdot oxy]}{methyl\}-2,3-dihydro[1,4]dioxino[2,3-c]pyridine,}$  or
  - a pharmaceutically acceptable salt thereof.
- 20. (Previously presented) A pharmaceutical composition, comprising a compound or salt according to claim 10 and a pharmaceutically acceptable carrier.
- 21. (Previously presented) A method of treating bacterial infections in a human, which comprises administering to a human in need thereof an effective amount of a compound or salt according to claim 10.
- 22. (New) A compound according to claim 1 wherein R<sup>10</sup> is H.